

# Health Update:

## Rabies Manufacturer's Recall of Human Rabies Vaccine

April 9, 2004

This document will be updated as new information becomes available. The current version can always be viewed at <http://www.dhss.state.mo.us/>.

The Missouri Department of Health & Senior Services (DHSS) is now using 4 types of documents to provide important information to medical and public health professionals, and to other interested persons:

**Health Alerts** convey information of the highest level of importance which warrants immediate action or attention from Missouri health providers, emergency responders, public health agencies, and/or the public.

**Health Advisories** provide important information for a specific incident or situation, including that impacting neighboring states; may not require immediate action.

**Health Guidances** contain comprehensive information pertaining to a particular disease or condition, and include recommendations, guidelines, etc. endorsed by DHSS.

**Health Updates** provide new or updated information on an incident or situation; can also provide information to update a previously sent Health Alert, Health Advisory, or Health Guidance; unlikely to require immediate action.

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April 9, 2004

FROM: RICHARD C. DUNN  
DIRECTOR  
SUBJECT: Manufacturer's Recall of Human Rabies Vaccine

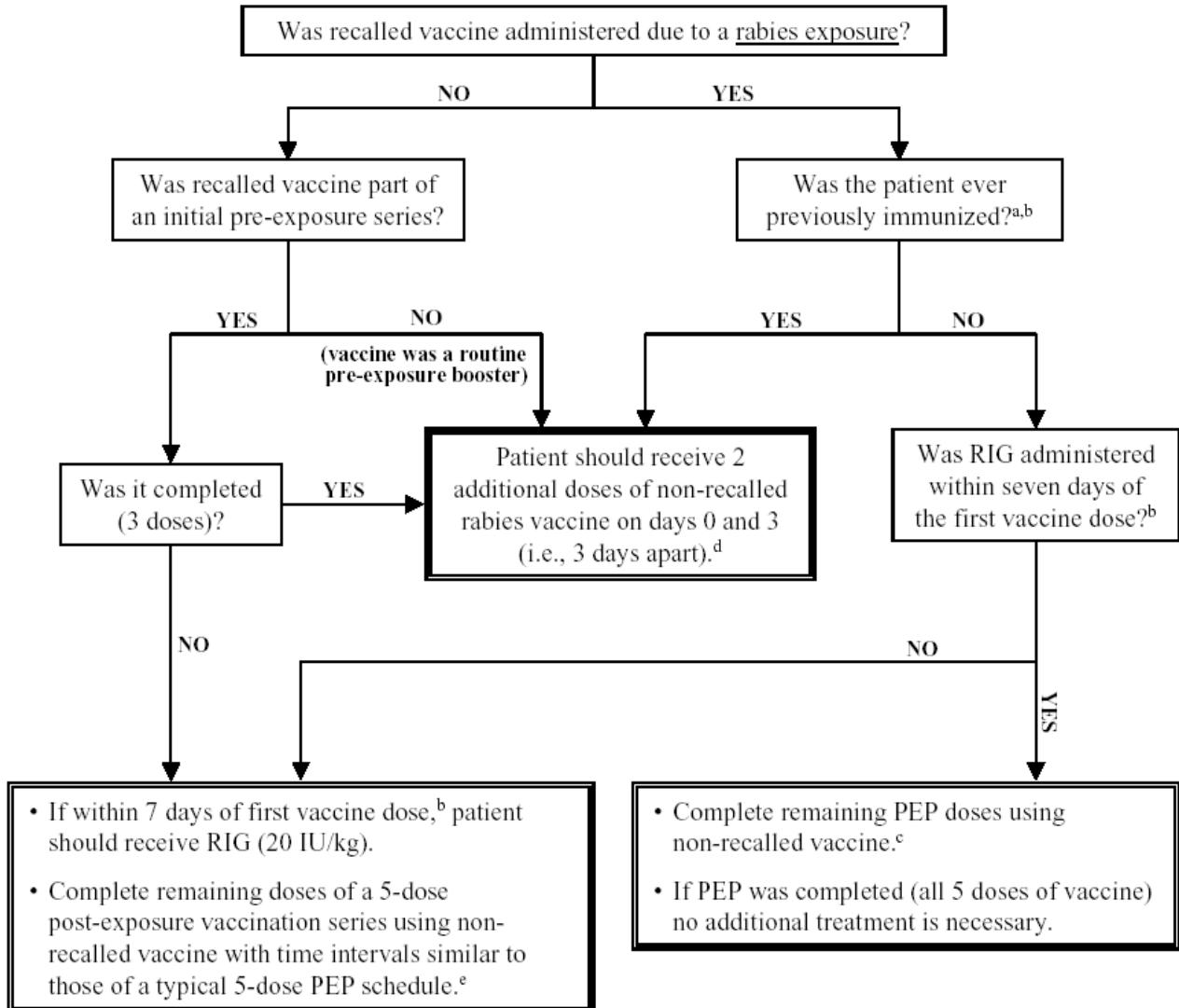
[This **Health Update** provides additional information on the manufacturer's recall of human rabies vaccine **Health Advisory** issued on April 3, 2004.]

The attached document is an algorithm for triage and disposition of persons who received Imovax rabies vaccine from 9/23/03 through 4/2/04 from vaccine lots X0667-2, X0667-3, W1419-2, and W1419-3. It was developed jointly by the New York State Health Department and the Centers for Disease Control and Prevention.

**Questions should be directed to the Department of Health and Senior Services 24 hours a day, 7 days a week at 800-392-0272.**

## Patient\* Triage, Imovax Rabies Vaccine Recall, April 2, 2004

(\*who received Imovax rabies vaccine from 9/23/03 through 4/2/04 from vaccine lots X0667-2, X0667-3, W1419-2, and W1419-3)



a. Previously immunized = 3 previous doses for pre-exposure or post-exposure.

b. Any vaccine regardless of whether that vaccine was in the recalled lots.

c. If PEP was discontinued because the animal was found to not have rabies, the patient should complete series using non-recalled vaccine (i.e., total of 5 doses of vaccine including any already given.)

d. If < 7 days since completion of 3 doses, the additional 2 doses should be given at days 7 and 21 after last previous dose.

e. Typical 5-dose PEP schedule = vaccine on days 0, 3, 7, 14, 28.